

REMARKS

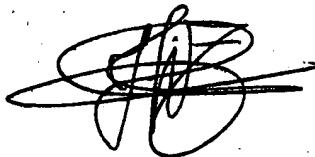
Claim 1 has been amended to introduce previously recited text that had been inadvertently omitted in amendment practice. The text introduced by this amendment is supported by the Application as filed. *See, e.g.*, Application, p. 2, *l.* 12, p. 6, *ll.* 17-18, p. 31, *l.* 20, p. 34, *l.* 17. The text introduced by this amendment was considered as such during examination until it was inadvertently omitted while editing the clean and marked-up versions of the claims in a previous amendment. *See, e.g.*, Amendment dated October 9, 2002, pp. 3, 5 (underlined text).

Applicants further submit that claim 1 as presently amended is comprised in claim Group I that was elected in response to the restriction requirement issued in the Office Action dated 03/26/2002, and that has been examined with the text introduced by the present amendment. *See, e.g.*, Amendment dated October 9, 2002, pp. 3, 5 (underlined text).

Applicants submit this Amendment and accompanying RCE for the purpose of clarifying the recitation of claim 1 and rendering its recitation as considered during examination. This Amendment corrects the inadvertent text omission that was made while editing text in a prior amendment, and it therefore makes it possible the issue of this Application as a U.S. patent with claim 1 printed therein free of the inadvertent text omission referred to hereinabove. In view of the above amendment and comments, Applicants respectfully submit that this Application is maintained in its condition for allowance, and passage to issue is earnestly requested.

Attached hereto is a marked-up version of the changes made to the claims by the current amendment. The attached page is captioned "Version with markings to show changes made."

Respectfully submitted,

A handwritten signature in black ink, appearing to be "J. Timoneda", with a large, stylized flourish extending to the left.

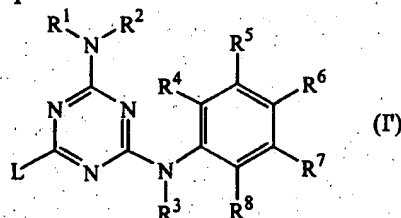
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Version with Markings to Show Changes Made

Amend Claim 1 as follows:

1. (Four times amended) A compound of formula



a pharmaceutically acceptable acid addition salt or a stereochemically isomeric form thereof, wherein

R¹ and R² are each independently selected from hydrogen; hydroxy; amino; C₁₋₆alkyl; C₁₋₆alkyloxy; C₁₋₆alkylcarbonyl; C₁₋₆alkyloxycarbonyl; Ar¹; mono- or di(C₁₋₆alkyl)amino; mono- or di(C₁₋₆alkyl)aminocarbonyl; dihydro-2(3*H*)-furanone; C₁₋₆alkyl substituted with one or two substituents each independently selected from amino, imino, aminocarbonyl, aminocarbonylamino, hydroxy, hydroxyC₁₋₆alkyloxy, carboxyl, mono- or di(C₁₋₆alkyl)amino, C₁₋₆alkyloxycarbonyl and thienyl; or

R¹ and R² taken together may form pyrrolidinyl, piperidinyl, morpholinyl, azido or mono- or di(C₁₋₆alkyl)aminoC₁₋₄alkylidene;

R³ is hydrogen, Ar¹, C₁₋₆alkylcarbonyl, C₁₋₆alkyl, C₁₋₆alkyloxycarbonyl, C₁₋₆alkyl substituted with C₁₋₆alkyloxycarbonyl; and

R⁴, R⁵, R⁷ and R⁸ are each independently selected from hydrogen, hydroxy, halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl or trihalomethyloxy;

R⁶ is aminocarbonyl; or

L is C₁₋₁₀alkyl; C₃₋₁₀alkenyl; C₃₋₁₀alkynyl; C₃₋₇cycloalkyl; or

L is C₁₋₁₀alkyl substituted with one or two substituents independently selected from the group consisting of C₃₋₇cycloalkyl; indolyl or indolyl substituted with one, two, three or four substituents each independently selected from halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethyloxy, or C₁₋₆alkylcarbonyl; and phenyl or phenyl substituted with one, two, three, four or five substituents each independently selected from halo, hydroxy, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethyloxy, or C₁₋₆alkylcarbonyl; and,

Ar¹ is phenyl, or phenyl substituted with one, two or three substituents each independently selected from halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, nitro or trifluoromethyl.